

Ph 203. Solution HW #2

1.)

Nuclear potential

(a) **Square well potential**

n-p triplet scattering:

$$a_t = -\lim_{k \rightarrow 0} \frac{\delta_0}{k},$$

where δ_0 is given by $kr_0 \left(\frac{\tan(kr_0)}{kr_0} - 1 \right)$, with $k = \sqrt{2mV_0}$.

n-p triplet bound state (deuteron):

$$K' \cot(K'r_0) = -k',$$

where $K' = \sqrt{2\mu(V_0 - E_B)}$ and $k' = \sqrt{2\mu E_B}$.

Need to solve numerically

$$\begin{cases} a_t = -r_0 \left(\frac{\tan(kr_0)}{kr_0} - 1 \right) \\ K' \cot(K'r_0) = -k', \end{cases}$$

for V_0, r_0 , given $a_t = 5.4$ fm, $E_B = 2.22$ MeV and $\mu \approx m_p/2 = 469$ MeV, which yields

$$r_0 \approx 2.1 \text{ fm}$$

$$V_0 \approx 34.2 \text{ MeV.}$$

(b) **Effective range r_e**

$$k \cot(\delta_0) = -\frac{1}{a_t} + \frac{r_e k^2}{2} + \dots$$

Need to find $\delta_0(k)$. Using the above equations and expanding,

$$\begin{aligned} \delta_0(k) &= -kr_0 + \tan^{-1}(-k(a_t - r_0)) \\ &= -kr_0 - k(a_t - r_0) - \frac{1}{3}k^3(a_t - r_0)^3 \\ &= -ka_t - \frac{1}{3}k^3(a_t - r_0)^3 + \dots \end{aligned}$$

Then we have

$$k \cot(\delta_0(k)) = -\frac{1}{a_t} + \frac{a_t^3 + (a_t - r_0)^3}{3a_t^3} k^2 + \mathcal{O}(k^3).$$

Therefore the effective range is given by (using numerical values from (a))

$$r_e = \frac{2}{3} \frac{a_t^3 + (a_t - r_0)^3}{a_t^2} \approx 4.4 \text{ fm.}$$

In Bertulani chapter 2.10, r_e is given for arbitrary potential using the binding energy $\rightarrow r_e \approx 1.76 \text{ fm}$. The radial square well potential, for which we computed r_e above, doesn't match the observed r_e very well $\rightarrow$ more complicated potential needed.

2.) Molecular deuterium ground state

a) molecular hydrogen: need overall wavefunction antisymmetric under $p \leftrightarrow p$. Let S be the total spin of pp and L the orbital angular momentum:

- (i) $S = 0$ (spin singlet, antisymmetric) $\rightarrow L = \text{even}$,
- (ii) $S = 1$ (spin triplet, symmetric) $\rightarrow L = \text{odd}$.

Assuming lower rotational modes have lower energy $\Rightarrow$ ground state $L = 0$ is a spin singlet (parahydrogen).

b) molecular deuterium: need wavefunction symmetric under $d \leftrightarrow d$ since deuteron has spin 1. Then we have

- (i) $S = 0$ (spin singlet, symmetric) $\rightarrow L = \text{even}$,
- (ii) $S = 1$ (spin triplet, antisymmetric) $\rightarrow L = \text{odd}$,
- (iii) $S = 2$ (5 states, symmetric) $\rightarrow L = \text{even}$.

$\Rightarrow$ ground state $L = 0$ is a symmetric spin state (ortho).

c) L is even for ortho, L odd for para

3.) Deuteron wavefunction

First, construct deuteron wavefunction for $j = 1, m_j = 1$ state

$$\psi(\vec{r}) = a\phi_{1s}(\vec{r}) + b\phi_{1d}(\vec{r}),$$

with the wavefunctions

$$\phi_{1s}(\vec{r}) = R_{1s}(r)Y_0^0(\theta, \phi) |1, 1\rangle,$$

$$\phi_{1d}(\vec{r}) = R_{1d}(r) \left(\sqrt{\frac{3}{5}} Y_2^2(\theta, \phi) |1, -1\rangle - \sqrt{\frac{3}{10}} Y_2^1(\theta, \phi) |1, 0\rangle + \sqrt{\frac{1}{10}} Y_2^0(\theta, \phi) |1, 1\rangle \right).$$

Radius: $r_0 \equiv \sqrt{\langle r^2 \rangle} = 1.96 \text{ fm}$

$$\begin{aligned} \langle r^2 \rangle &= \langle \psi | r^2 | \psi \rangle \\ &= |a|^2 \int_0^\infty dr r^4 |R_{1s}|^2 + |b|^2 \int_0^\infty dr r^4 |R_{1d}|^2 \\ &= |a|^2 \frac{3}{2\alpha} + |b|^2 \frac{7}{2\alpha} = \frac{3}{2\alpha} + |b|^2 \frac{2}{\alpha}, \end{aligned}$$

using $|a|^2 = 1 - |b|^2$. We assume $|b| \ll 1 \Rightarrow$ to linear order in b , we have

$$\langle r^2 \rangle \cong \frac{3}{2\alpha}.$$

Quadrupole moment: $\hat{Q} = e \sqrt{\frac{16\pi}{5}} r^2 Y_2^0(\theta, \phi)$, $q_0 \equiv \langle \hat{Q} \rangle = 0.286 \text{ efm}$

$$\begin{aligned} \langle \hat{Q} \rangle &= |a|^2 \int d^3r \phi_{1s}^* \hat{Q} \phi_{1s} + |b|^2 \int d^3r \phi_{1d}^* \hat{Q} \phi_{1d} \\ &\quad + 2\text{Re}[ab^* \int d^3r \phi_{1d}^* \hat{Q} \phi_{1s}], \end{aligned}$$

$\rightarrow$ first term vanishes. This is due to the Y_{lm} orthogonality (i.e $\langle Y_{00} | Y_{20} \rangle = 0$)

$\rightarrow$ second term can be neglected to linear order in $|b|$

$\rightarrow$ third term: only non-vanishing term is spin $|1, 1\rangle$ term

$$\int d^3r \phi_{1d}^* \hat{Q} \phi_{1s} = e \sqrt{\frac{16\pi}{5}} \sqrt{\frac{1}{10}} \int dr r^4 R_{1s} R_{1d} \int d\Omega Y_2^{0*} Y_2^0 Y_0^0 = \frac{e}{2\alpha} \sqrt{\frac{6}{5}}$$

$$\Rightarrow \langle \hat{Q} \rangle = \frac{e}{\alpha} \sqrt{\frac{6}{5}} \text{Re}[b] = \frac{2er_0^2}{3} \sqrt{\frac{6}{5}} \text{Re}[b]$$

$$\Rightarrow \text{for } b \in \mathbf{R}, b \approx \frac{3q_0}{2er_0^2} \sqrt{\frac{5}{6}} \approx 0.1,$$

so deuteron has approximately 1% d -wave and 99% s -wave.

4.)

Scattering rate for spin flip

Scattering amplitude $f(\theta) \propto \langle f | \hat{H} | i \rangle$, where $|i\rangle$ and $|f\rangle$ denote the initial and final state, respectively. For low energy scattering $k \rightarrow 0$ (s -wave scattering only):

$$f(\theta) \approx -a,$$

where a is the scattering length.

Total scattering rate: average over initial states, sum over final states

$$\begin{aligned} \sigma_{tot} &\propto \frac{1}{4} \sum_{i,f} |\langle f | \hat{H} | i \rangle|^2 \\ &\propto \frac{1}{4} \left(|\langle ++ | \hat{H} | ++ \rangle|^2 + |\langle ++ | \hat{H} | +- \rangle|^2 + |\langle ++ | \hat{H} | -+ \rangle|^2 + \dots \right) \\ &\propto \frac{1}{4} \left(|\langle ++ | \hat{H} | ++ \rangle|^2 + |\langle +- | \hat{H} | +- \rangle|^2 + |\langle +- | \hat{H} | -+ \rangle|^2 \right. \\ &\quad \left. + |\langle -+ | \hat{H} | +- \rangle|^2 + |\langle -+ | \hat{H} | -+ \rangle|^2 + |\langle -- | \hat{H} | -- \rangle|^2 \right), \end{aligned}$$

where in the last line we wrote down only the non-zero matrix elements.

Recall that

$$\begin{aligned} |+- \rangle &= \frac{1}{\sqrt{2}} (|1, 0 \rangle + |0, 0 \rangle) \\ |-+ \rangle &= \frac{1}{\sqrt{2}} (|1, 0 \rangle - |0, 0 \rangle). \end{aligned}$$

Hence we obtain

$$\begin{aligned} \langle +- | \hat{H} | +- \rangle &= \frac{1}{2} (a_t + a_s), \\ \langle +- | \hat{H} | -+ \rangle &= \frac{1}{2} (a_t - a_s), \\ \langle -+ | \hat{H} | +- \rangle &= \frac{1}{2} (a_t - a_s), \\ \langle -+ | \hat{H} | -+ \rangle &= \frac{1}{2} (a_t + a_s). \end{aligned}$$

Plugging back to the expression for the total cross-section

$$\begin{aligned}\sigma_{tot} &\propto \frac{1}{4} \left(a_t^2 + 2 \times \frac{1}{4} (a_t + a_s)^2 + 2 \times \frac{1}{4} (a_t - a_s)^2 + a_t^2 \right) \\ &\propto \frac{3}{4} a_t^2 + \frac{1}{4} a_s^2.\end{aligned}$$

Repeating the above steps for the cross-section for a spin flip

$$\begin{aligned}\sigma_{flip} &\propto \frac{1}{4} \left(\left| \langle + - | \hat{H} | - + \rangle \right|^2 + \left| \langle - + | \hat{H} | + - \rangle \right|^2 \right) \\ &\propto \frac{1}{4} \left(2 \times \frac{1}{4} (a_t - a_s)^2 \right) \\ &\propto \frac{1}{8} (a_t - a_s)^2.\end{aligned}$$

Therefore the probability of spin flip is

$$\frac{\sigma_{flip}}{\sigma_{tot}} = \frac{(a_t - a_s)^2}{2(3a_t^2 + a_s^2)} \approx 65\%.$$

5.) Bertulani 2.4 and 2.12

a) potential well with hard core

$$V(r) = \begin{cases} \infty & \text{for } r < c \\ -V_0 & \text{for } c < r < R + c \\ 0 & \text{for } r > R + c \end{cases}$$

Radial part of bound state wavefunction (following Bertulani):

$$u(r) = \begin{cases} 0 & \text{for } r < c \\ A \sin [K(r - c)] & \text{for } c < r < R + c \\ B e^{-kr} & \text{for } r > R + c, \end{cases}$$

where $k = \sqrt{2\mu E_B}$ and $K = \sqrt{2\mu(V_0 - E_B)}$.

Boundary matching at $r = R + c$:

$$\text{continuity: } A \sin [KR] = B e^{-k(R+c)}$$

$$\text{1st derivative: } AK \cos [KR] = -Bke^{-k(R+c)}$$

$$\Rightarrow K \cot [KR] = -k.$$

b) scattering off a hard sphere potential

$$V(r) = \begin{cases} \infty & \text{for } r < r_0 \\ 0 & \text{for } r > r_0 \end{cases}$$

Radial wavefunction

$$u(r) = \begin{cases} 0 & \text{for } r < r_0 \\ A \sin (kr + \delta_0) & \text{for } r > r_0. \end{cases}$$

Boundary matching at $r = r_0$: $A \sin (kr_0 + \delta_0) = 0$

$$\Rightarrow \delta_0 = -kr_0.$$

Hence the cross-section for low energy scattering ($k \ll \frac{1}{r_0}$):

$$\sigma = \frac{4\pi}{k^2} \sin^2 \delta_0 \approx \frac{4\pi}{k^2} \delta_0^2 = 4\pi r_0^2.$$